

What is claimed is:

1. An isolated polypeptide that stimulates gastrointestinal smooth muscle contraction, comprising an amino acid sequence at least 80% identical to the
5 sequence of human prokineticin 1 (SEQ ID NO:3), said sequence comprising the N-terminal 6 amino acids of SEQ ID NO:3, the 10 conserved cysteine residues of SEQ ID NO:3, and from 0 to 9 of the 9 C-terminal amino acids of SEQ ID NO:3.
- 10 2. The isolated polypeptide of claim 1, wherein amino acid residues that differ from residues in SEQ ID NO:3 are conservative substitutions thereof.
3. The isolated polypeptide of claim 1, wherein amino acid residues that differ from residues in
15 SEQ ID NO:3 consist of the corresponding residues from SEQ ID NO:6.
4. The isolated polypeptide of claim 3, comprising SEQ ID NO:13.
5. The isolated polypeptide of claim 1,
20 comprising amino acids 1-77 of SEQ ID NO:3.
6. The isolated polypeptide of claim 1, comprising SEQ ID NO:3.
7. The isolated polypeptide of claim 1, comprising a 6XHis tag.

8. The isolated polypeptide of claim 1, which is detectably labeled.

9. An isolated peptide comprising at least 10 contiguous amino acids of SEQ ID NO:3, wherein said peptide is immunogenic.

10. A pharmaceutical composition, comprising the isolated polypeptide of claim 1 and a pharmaceutically acceptable carrier.

11. A method of stimulating gastrointestinal smooth muscle contraction in a mammal, comprising administering to said mammal an effective amount of the polypeptide of claim 1.

12. A nucleic acid molecule encoding the polypeptide of claim 1.

13. An expression vector containing the nucleic acid molecule of claim 12 operatively linked to a promoter of gene expression.

14. A host cell comprising the expression vector of claim 13.

15. A method of preparing the isolated polypeptide of claim 1, comprising culturing the host cell of claim 14 so as to express said polypeptide, substantially purifying said polypeptide, and refolding said polypeptide.

16. An antibody that selectively binds the polypeptide of claim 1.

17. An isolated polypeptide that stimulates gastrointestinal smooth muscle contraction, comprising an amino acid sequence at least 80% identical to the sequence of human prokineticin 2 (SEQ ID NO:6), said sequence comprising the N-terminal 6 amino acids of SEQ ID NO:6, the 10 conserved cysteine residues of SEQ ID NO:6, and from 0 to 4 of the 4 C-terminal amino acids of SEQ ID NO:6.

18. The isolated polypeptide of claim 17, wherein amino acid residues that differ from residues in SEQ ID NO:6 are conservative substitutions thereof.

19. The isolated polypeptide of claim 17, wherein amino acid residues that differ from residues in SEQ ID NO:6 consist of the corresponding residues from SEQ ID NO:3.

20. The isolated polypeptide of claim 19, comprising SEQ ID NO:14.

21. The isolated polypeptide of claim 17, comprising amino acids 1-77 of SEQ ID NO:6.

22. The isolated polypeptide of claim 17, comprising SEQ ID NO:6.

23. The isolated polypeptide of claim 17, comprising a 6XHis tag.

24. The isolated polypeptide of claim 17,
which is detectably labeled.

25. An isolated peptide comprising at least 10
contiguous amino acids of SEQ ID NO:6, wherein said
5 peptide is immunogenic.

26. A pharmaceutical composition, comprising
the isolated polypeptide of claim 17 and a
pharmaceutically acceptable carrier.

27. A method of stimulating gastrointestinal
10 smooth muscle contraction in a mammal, comprising
administering to said mammal an effective amount of the
polypeptide of claim 17.

28. A nucleic acid molecule encoding the
polypeptide of claim 17.

29. An expression vector containing the
15 nucleic acid molecule of claim 17 operatively linked to a
promoter of gene expression.

30. A host cell comprising the expression
vector of claim 29.

31. A method of preparing the isolated
20 polypeptide of claim 17, comprising culturing the host
cell of claim 30 so as to express said polypeptide,
substantially purifying said polypeptide, and refolding
said polypeptide.

32. An antibody that selectively binds the polypeptide of claim 17.

33. A method of identifying a prokineticin receptor ligand, comprising contacting a preparation
5 comprising prokineticin receptor with one or more candidate compounds, and identifying a compound that specifically binds to said receptor, said compound being characterized as a prokineticin receptor ligand.

34. The method of claim 33, wherein said
10 preparation is an intestinal smooth muscle preparation or membrane preparation thereof.

35. The method of claim 33, wherein said preparation is a cell line or membrane preparation thereof.

36. The method of claim 35, wherein said cell
15 line is M2A7 (ATCC CRL-2500).

37. The method of claim 33, wherein the ability of said ligand to selectively agonize or antagonize prokineticin receptor signaling is further
20 determined.

38. The method of claim 37, wherein said signaling is determined in a cell line.

39. The method of claim 38, wherein said cell line is M2A7 (ATCC CRL-2500).

40. The method of claim 37, wherein said signaling is determined by monitoring calcium mobilization.

41. The method of claim 33, wherein the
5 ability of said ligand to modulate smooth muscle contractility is further determined.

42. A method of identifying a prokineticin
receptor agonist, comprising contacting a preparation
comprising a prokineticin receptor with one or more
10 candidate compounds, and identifying a compound that
selectively promotes production of a prokineticin
receptor signal, said compound being characterized as a
prokineticin receptor agonist.

43. The method of claim 42, wherein said
15 preparation is a cell line.

44. The method of claim 43, wherein said cell
line is M2A7 (ATCC CRL-2500).

45. The method of claim 42, wherein said
signaling is determined by monitoring calcium
20 mobilization.

46. The method of claim 42, wherein the
ability of said agonist to modulate smooth muscle
contractility is further determined.

47. A method of identifying a prokineticin
25 receptor antagonist, comprising contacting a preparation
comprising a prokineticin receptor with one or more

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5 antagonist.

NOS:3 and amino acids 1-77 of SEQ ID NO:6.

10 49. The method o
preparation is a cell line.

50. The method of claim 49, wherein said cell line is M2A7 (ATCC CRL-2500).

15 signaling is determined by monitoring calcium
mobilization.

52. The method of claim 47, wherein the ability of said antagonist to modulate smooth muscle contractility is further determined.